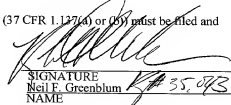


60 Rec'd PCT/PTO 20 JUL 2001

Form PTO-1390		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER P21259
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		U.S. APPLICATION NO. (If known, see 37 CFR 1.5) 09/869310	
INTERNATIONAL APPLICATION NO. PCT/AT00/00102	INTERNATIONAL FILING DATE 20 April 2000	PRIORITY DATE CLAIMED 22 April 1999	
TITLE OF INVENTION CASTING MATERIAL FOR INDEFINITE ROLLERS WITH A SLEEVE PART AND METHOD FOR PRODUCING THE SAME			
APPLICANT(S) FOR DO/EO/US Bernhard FEISTRITZER, Karl-Heinrich SCHRÖDER, Michael WINDHAGER, and Karl-Heinz ZIEHENBERGER			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information.			
<ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. 3. <input checked="" type="checkbox"/> This is an express request to promptly begin national examination procedures (35 U.S.C. 371(f)). 4. <input checked="" type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (PCT Article 31). 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)). <ol style="list-style-type: none"> a. <input checked="" type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> has been communicated by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). 6. <input type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)). 7. <input type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> have been communicated by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input type="checkbox"/> have not been made and will not be made. 8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)) 9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). 10. <input type="checkbox"/> "Unexecuted" 10. <input type="checkbox"/> An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (U.S.C. 371(c)(5)). 			
Items 11 to 16 below concern other document(s) or information included:			
11. Assignee: <u>EISENWERK SULZAU-WERFEN R&E WEINBERGER AG of Tenneck, AUSTRIA</u>			
12. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.			
13. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.			
14. <input type="checkbox"/> A FIRST preliminary amendment.			
15. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.			
16. <input type="checkbox"/> A substitute specification.			
17. <input type="checkbox"/> A change of power of attorney and/or address letter.			
18. <input type="checkbox"/> Figure of Drawing to be published			
19. <input checked="" type="checkbox"/> Other items or information: Cover Sheet and International Application as published in German. Two PCT/PEA/408 Forms (in German). Response to First Written Opinion (in German). Cover Letter under 35 USC 371 and 1.495. Claim of Priority.			

U.S. APPLICATION NO. (if known, see 37 CFR 1.5) 09/869310	INTERNATIONAL APPLICATION NO. PCT/AT00/00102	ATTORNEY'S DOCKET NUMBER P21259	
19. The following fees are submitted: Basic National Fee (37 CFR 1.492(a)(1)-(5)): Search report has been prepared by the EPO or JPO. \$ 860.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) \$ 690.00 No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)) \$ 710.00 Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO. \$1,000.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4). \$ 100.00 ENTER APPROPRIATE BASIC FEE AMOUNT =		CALCULATIONS	PTO USE ONLY
Surcharge of \$130.00 for furnishing the oath or declaration later than ___ 20 ___ 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		\$	
Claims	Number Filed	Number Extra	RATE
Total Claims	- 20 =		X \$18.00 \$
Independent Claims	- 3 =		X \$80.00 \$
Multiple dependent claim(s) (if applicable)			+ \$270.00 \$
TOTAL OF ABOVE CALCULATIONS =			\$860.00
Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by ____%.			\$
SUBTOTAL =			\$860.00
Processing fee of \$130.00 for furnishing the English translation later than ___ 20 ___ 30 months from the earliest claimed priority date (37 CFR 1.492(f)).			+
Extension of Time fee in the amount of \$			
TOTAL NATIONAL FEE =			\$860.00
Fee for recording the enclosed assignment (37 CFR 1.21(h). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property			+
TOTAL FEES ENCLOSED =			\$860.00
			Amount to be refunded \$
			Charged \$
a. <input checked="" type="checkbox"/> A check in the amount of <u>\$860.00</u> to cover the above fees is enclosed. b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>19-0089</u> . NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.127(d) or (e)) must be filed and granted to restore the application to pending status. SEND ALL CORRESPONDENCE TO CUSTOMER NO. 7055 AT THE PRESENT ADDRESS OF: Neil F. Greenblum GREENBLUM & BERNSTEIN, P.L.C. 1941 Roland Clarke Place Reston, VA 20191 (703) 716-1191			
 SIGNATURE Neil F. Greenblum NAME			<u>35,973</u> 28,394 REGISTRATION NUMBER

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant : Bernhard FEISTRITZER

Group Art Unit: Unknown

Serial No : 09/869,310

Examiner: Unknown

Filed : July 20, 2001

For : CASTING MATERIAL FOR INDEFINITE ROLLERS WITH A
SLEEVE PART AND METHOD FOR PRODUCING THE SAME**PRELIMINARY AMENDMENT**Commissioner of Patents and Trademarks
Washington, DC 20231

Sir :

Prior to examination of the application and calculation of filing fees, please enter the following amendment.

IN THE SPECIFICATION

Please add the Abstract appended as Appendix 1 on a separate page at the end of this Amendment.

IN THE CLAIMS

Please amend claims 3-12, 15-25, and 28-30 as follows (for the Examiner's convenience, a clean copy of all pending claims is being reproduced below, with amended claims 3-12, 15-25, and 28-30 being so labeled, and a marked-up version of the amended

claims being submitted in Appendix 2 attached at the end of this Amendment):

CLAIMS

1. Method for the production and processing of alloyed casting material for the working area of indefinite chill rolls, containing the elements carbon, silicon, manganese, chromium, nickel, molybdenum, vanadium, and if applicable additional elements of group 5 of the periodic system, aluminum, and the remainder iron, accompanying elements, and impurities related to the manufacturing process, characterized in that
 - A. a melt is produced that has a chemical composition in wt-% of
 - 2.0 to 3.5 C
 - 1.0 to 2.0 Si
 - 0.5 to 2.0 Mn
 - 1.0 to 3.0 Cr
 - 3.5 to 4.9 Ni
 - 0.2 to 2.9 Mowith the remainder iron and impurities, and
 - B. more than 0.5 % vanadium by weight in amounts up to 5.9 wt-% is added, is dissolved therein, and
 - C. the composition of the melt is set using alloying methods by fixing the

concentrations of carbon and silicon in the presence of nickel and the effective total of the carbide forming elements in such a manner that, at its solidification, a microstructure is formed which exhibits 1.0 to 3.0 vol-% of graphite, with the guideline that more than 20 but less than 100 graphite particles are present per mm² of observed surface in a metallographic section and the remainder is composed primarily of martensite, 8 to 35 vol-% of eutectic carbides, and at least 1 vol-% of finely distributed vanadium carbides, after which

D. the melt is cast in a form, preferably a centrifugal casting mold, and is allowed to solidify into a body, preferably a working body of a roll, and optionally, the cast body is further processed, for example, into a composite roll, with the body or roll that has been produced in this manner being

E. subjected to a heat treatment including at least one one-time heating to a treatment temperature, holding at this temperature, and cooling to room temperature.

2. Method according to claim 1, characterized in that the composition of the melt is set using alloying methods by fixing the concentrations of carbon and silicon in the presence of nickel and the effective total of the carbide forming elements in such a manner that, at its solidification, a microstructure is formed which has 1.0 to 2.5 vol-% of graphite, with the guideline that more than 22 and less than 100 graphite

particles are present per mm² of observed surface in a metallographic section and the remainder is composed primarily of martensite, 10 to 25 vol-% of eutectic carbides, and of 2 to 20% of finely distributed carbides of the elements of group 5 of the periodic system.

3. (Amended) Method according to claim 1, characterized in that the composition of the melt is set in such a manner that the concentration ratio of carbon to silicon is less than or equal to 2.6, preferably less than or equal to 2.0:

$$C/Si \leq 2.6, \text{ preferably } = 2.0$$

4. (Amended) Method according to claim 1, characterized in that the carbon content of the melt is set to a value of 2.2 to 3.1 wt-%, preferably 2.6 to 2.95%.

5. (Amended) Method according to claim 1, characterized in that a final content of silicon of 1.2 to 1.85 wt-%, preferably 1.4 to 1.75%, is provided.

6. (Amended) Method according to claim 1, characterized in that, when the composition of the melt is set using alloying methods, 0.002 to 0.65 wt-%, preferably 0.005 to 0.04

%, of aluminum is added and dissolved therein.

7. (Amended) Method according to claim 1, characterized in that the nickel content of the melt is set to a value of 3.51 to 4.7 wt-%, preferably 4.15 to 4.6 wt-%.

8. (Amended) Method according to claim 1, characterized in that the composition of the melt is set in such a manner that the concentration ratio of molybdenum to chromium is less than 1.0, preferably less than 0.8:

$\text{Mo/Cr} < 1.0$, preferably < 0.8 .

9. (Amended) Method according to claim 1, characterized in that the content levels of chromium and molybdenum in wt-% of the melt are set to the values of
chromium 1.5 to 1.9
molybdenum 0.3 to 0.9.

10. (Amended) Method according to claim 1, characterized in that 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.9 wt-%, is added to the melt and dissolved therein.

11. (Amended) Method according to claim 1, characterized in that some of the vanadium is replaced by additional elements from group 5 of the periodic system in an amount of less than 0.6 wt-%, and mixed carbides are formed.
12. (Amended) Method according to claim 1, characterized in that the cast body or the roll is subjected to a heat treatment which comprises heating from room temperature to a treatment temperature of 400 °C to 500 °C, preferably 460 °C to 480 °C, holding at this temperature for at least two hours, preferably at least 8 hours, and cooling to room temperature, optionally with a low-temperature treatment.
13. Casting material for the working area of indefinite chill rolls, containing the elements carbon, silicon, manganese, chromium, nickel, molybdenum, vanadium, and the remainder iron, accompanying elements, and impurities related to the manufacturing process, characterized in that the alloy contains, in wt-%, more than
 - 0.5 to 5.9 vanadium
 - 1.0 to 2.0 silicon
 - 0.5 to 2.0 manganese
 - 1.0 to 3.0 chromium
 - 3.5 to 4.9 nickel

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0.20 to 2.9 molybdenum

2.0 to 3.5 carbon with the guideline that

1.0 to 3.0 vol-% is present as graphite in particles with a distribution of more than 20 and less than 100 particles per mm² of polished surface of the material.

14. Casting material according to claim 13, characterized in that the alloy contains

1.8 to 4.9 wt-% vanadium

2.2 to 3.1 wt-% carbon with the guideline that

1.2 to 2.5 vol-% as graphite in particles with a distribution of more than 22 and less than 90 particles per mm² of polished surface.

15. (Amended) Casting material according to claim 13, characterized in that the alloy contains, in wt-%,

2.0 to 3.5 carbon

1.0 to 2.0 silicon

0.5 to 2.0 manganese

1.2 to 2.5 chromium

3.5 to 4.9 nickel

0.5 to 2.1 molybdenum

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1.5 to 4.9 vanadium

with the remainder iron and impurities.

16. (Amended) Casting material according to claim 13, characterized in that the alloy has a concentration ratio of carbon to silicon of less than or equal to 2.6, preferably less than or equal to 2.0:

$C/Si \leq 2.6$, preferably ≤ 2.0

17. (Amended) Casting material according to claim 13, characterized in that the alloy contains 2.6 to 2.95% carbon by weight.
18. (Amended) Casting material according to claim 13, characterized in that the alloy contains 1.2 to 1.85 wt-% of silicon, preferably 1.4 to 1.75%
19. (Amended) Casting material according to claim 13, characterized in that the alloy contains 0.002 to 0.65 wt-% of aluminum, preferably 0.005 to 0.04 %.
20. (Amended) Casting material according to claim 13, characterized in that the alloy

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contains 3.5 to 4.9 wt-% of nickel, preferably 4.15 to 4.6%.

21. (Amended) Casting material according to claim 13, characterized in that the alloy has a concentration ratio of molybdenum to chromium of less than 1.0, preferably less than 0.8:

$\text{Mo/Cr} < 1.0$, preferably < 0.8 .

22. (Amended) Casting material according claim 13, characterized in that the alloy contains in wt-%
chromium 1.5 to 2.01
molybdenum 0.3 to 0.9.

23. (Amended) Casting material according to claim 13, characterized in that the alloy contains 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.95 wt-%.

24. (Amended) Casting material according to claim 13, characterized in that some of the vanadium content is replaced by additional elements from group 5 of the periodic system in a proportion of less than 0.6 wt-%.

25. (Amended) Casting material according to claim 13, characterized in that the material possesses, in vol-%,
- 8 to 35, preferably 10 to 25, eutectic carbides,
- and 1 to 15, preferably 2 to 10, carbides of the elements of group 5 of the periodic system.
26. Composite indefinite chill rolls, especially for finishing stands of wide strip rolling mills and also Steckel and heavy plate mills, preferably produced using a method according to claims 1 through 12, comprising a work or sleeve part made of a casting alloy, preferably according to claims 13 through 25, with little tendency to adhere or weld to the rolling stock and with a high-strength core part made of ductile iron, characterized in that the working area or sleeve has a thickness of 10 to 150 mm and the sleeve material a structure composed essentially of 1.0 to 2.5 vol-% of graphite, with the latter being finely dispersed with a graphite particle count of more than 20 particles per mm² in a metallographic section, and of 8 to 35 vol-% of eutectic carbides, and of 1 to 20 vol-% of uniformly distributed vanadium carbide, with the remainder composed primarily of martensite and constituents related to impurities or the manufacturing process, and has a hardness between 70 and 90 ShC.

27. Composite indefinite chill roll according to claim 26, characterized in that the working area or sleeve material has a structure which has 1.0 to 2.5 vol-% of graphite, with the guideline that its distribution density is at least 22 particles but less than 100 graphite particles per mm² of polished surface in a metallographic section, contains eutectic carbides in an amount of 10 to 25 vol-%, and 2 to 10 vol-% special carbides of the elements of group 5 of the periodic system.

28. (Amended) Composite indefinite chill rolls according to claim 26, characterized in that the working or sleeve material has a composition, in wt-%, of

C = 2.0 to 3.5, preferably 2.21 to 3.1, especially 2.6 to 2.95

Si = 1.0 to 2.0, preferably over 1.2 to 1.85, especially 1.4 to 1.75

Mn = 0.5 to 2.0, preferably 0.6 to 1.6, especially 0.7 to 1.4

Cr = 1.0 to 3.0, especially 1.5 to 2.01

Ni = 3.5 to 4.9, preferably 3.5 to 4.7, especially 4.15 to 4.6

Mo = 0.20 to 2.9, especially 0.3 to 0.9

Al = 0.002 to 0.65, preferably 0.005 to 0.1, especially 0.005 to 0.04

V = 0.5 to 5.9, preferably 1.8 to 3.9, especially 1.9 to 2.9

with the remainder being iron and impurities and the roll core is formed of ductile iron.

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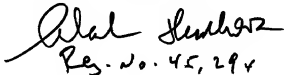
29. (Amended) Composite indefinite chill rolls according to claim 26, characterized in that the working or sleeve material has, in wt-%,
- V = 3.1 to 3.9, preferably 3.3 to 3.75 and
- Nb+Ta = less than 0.6
- with the remainder being iron and impurities.
30. (Amended) Composite indefinite chill rolls according to claim 26, characterized in that the binding zone between the sleeve or working part and the roll core of low-alloy cast iron, preferably of ductile iron, has, in the radial direction, a bending strength (3-point bending test) of greater than 600 N/mm².

Remarks

The changes above were made simply to remove multiple dependency in the claims and provide an Abstract of the Disclosure as required by the regulations.

Should the Examiner have any further comments or questions, the Examiner is invited to contact the undersigned at the below-listed telephone number.

Respectfully submitted,
Bernhard FEISTRITZER



Reg. No. 45,294

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Enclosure: Appendices 1 and 2

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a composite roll, with the body or roll that has been produced in this manner being subjected to a heat treatment including at least one one-time heating to a treatment temperature, holding at this temperature, and cooling to room temperature.--

Appendix 2

Claims

3. (Amended) Method according to [one of claims 1 or 2] claim 1, characterized in that the composition of the melt is set in such a manner that the concentration ratio of carbon to silicon is less than or equal to 2.6, preferably less than or equal to 2.0:

$$\text{C/Si} \leq 2.6, \text{ preferably } = 2.0$$

4. (Amended) Method according to [one of claims 1 through 3] claim 1, characterized in that the carbon content of the melt is set to a value of 2.2 to 3.1 wt-%, preferably 2.6 to 2.95%.
5. (Amended) Method according to [one of claims 1 through 4] claim 1, characterized in that a final content of silicon of 1.2 to 1.85 wt-%, preferably 1.4 to 1.75%, is provided.
6. (Amended) Method according to [one of claims 1 through 5] claim 1, characterized in that, when the composition of the melt is set using alloying methods, 0.002 to 0.65 wt-%, preferably 0.005 to 0.04 %, of aluminum is added and dissolved therein.
7. (Amended) Method according to [one of claims 1 through 7] claim 1, characterized in that the nickel content of the melt is set to a value of 3.51 to 4.7 wt-%, preferably 4.15 to 4.6 wt-%.

%.

8. (Amended) Method according to [one of claims 1 through 7] claim 1, characterized in that the composition of the melt is set in such a manner that the concentration ratio of molybdenum to chromium is less than 1.0, preferably less than 0.8:

Mo/Cr < 1.0, preferably < 0.8.

9. (Amended) Method according to [one of claims 1 through 8] claim 1, characterized in that the content levels of chromium and molybdenum in wt-% of the melt are set to the values of
- chromium 1.5 to 1.9
- molybdenum 0.3 to 0.9.

10. (Amended) Method according to [one of claims 1 through 9] claim 1, characterized in that 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.9 wt-%, is added to the melt and dissolved therein.

11. (Amended) Method according to [one of claims 1 through 10] claim 1, characterized in that some of the vanadium is replaced by additional elements from group 5 of the periodic system in an amount of less than 0.6 wt-%, and mixed carbides are formed.

12. (Amended) Method according to [one of claims 1 through 11] claim 1, characterized in that the cast body or the roll is subjected to a heat treatment which comprises heating from room temperature to a treatment temperature of 400 °C to 500°C, preferably 460 °C to 480 °C, holding at this temperature for at least two hours, preferably at least 8 hours, and cooling to room temperature, optionally with a low-temperature treatment.
15. (Amended) Casting material according to claim 13 [or 14], characterized in that the alloy contains, in wt-%,
- | | |
|------------|------------|
| 2.0 to 3.5 | carbon |
| 1.0 to 2.0 | silicon |
| 0.5 to 2.0 | manganese |
| 1.2 to 2.5 | chromium |
| 3.5 to 4.9 | nickel |
| 0.5 to 2.1 | molybdenum |
| 1.5 to 4.9 | vanadium |
- with the remainder iron and impurities.
16. (Amended) Casting material according to [one of claims 13 through 15] claim 13, characterized in that the alloy has a concentration ratio of carbon to silicon of less than or equal to 2.6, preferably less than or equal to 2.0:

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$C/Si \leq 2.6$, preferably ≤ 2.0

17. (Amended) Casting material according to [one of claims 13 through 16] claim 13, characterized in that the alloy contains 2.6 to 2.95% carbon by weight.
18. (Amended) Casting material according to [one of claims 13 through 17] claim 13, characterized in that the alloy contains 1.2 to 1.85 wt-% of silicon, preferably 1.4 to 1.75%
19. (Amended) Casting material according to [one of claims 13 through 18] claim 13, characterized in that the alloy contains 0.002 to 0.65 wt-% of aluminum, preferably 0.005 to 0.04 %.
20. (Amended) Casting material according to [one of claims 13 through 19] claim 13, characterized in that the alloy contains 3.5 to 4.9 wt-% of nickel, preferably 4.15 to 4.6%.
21. (Amended) Casting material according to [one of claims 13 through 20] claim 13, characterized in that the alloy has a concentration ratio of molybdenum to chromium of less than 1.0, preferably less than 0.8:

$Mo/Cr < 1.0$, preferably < 0.8 .

22. (Amended) Casting material according [to one of claims 13 through 21] claim 13, characterized in that the alloy contains in wt-%
chromium 1.5 to 2.01
molybdenum 0.3 to 0.9.
23. (Amended) Casting material according to [one of claims 13 through 22] claim 13, characterized in that the alloy contains 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.95 wt-%.
24. (Amended) Casting material according to [one of claims 13 through 23] claim 13, characterized in that some of the vanadium content is replaced by additional elements from group 5 of the periodic system in a proportion of less than 0.6 wt-%.
25. (Amended) Casting material according to [one of claims 13 through 24] claim 13, characterized in that the material possesses, in vol-%,
8 to 35, preferably 10 to 25, eutectic carbides,
and 1 to 15, preferably 2 to 10, carbides of the elements of group 5 of the periodic system.
28. (Amended) Composite indefinite chill rolls according to claim 26 [or 27], characterized in that the working or sleeve material has a composition, in wt-%, of
C = 2.0 to 3.5, preferably 2.21 to 3.1, especially 2.6 to 2.95

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- Si = 1.0 to 2.0, preferably over 1.2 to 1.85, especially 1.4 to 1.75
- Mn = 0.5 to 2.0, preferably 0.6 to 1.6, especially 0.7 to 1.4
- Cr = 1.0 to 3.0, especially 1.5 to 2.01
- Ni = 3.5 to 4.9, preferably 3.5 to 4.7, especially 4.15 to 4.6
- Mo = 0.20 to 2.9, especially 0.3 to 0.9
- Al = 0.002 to 0.65, preferably 0.005 to 0.1, especially 0.005 to 0.04
- V = 0.5 to 5.9, preferably 1.8 to 3.9, especially 1.9 to 2.9

with the remainder being iron and impurities and the roll core is formed of ductile iron.

29. (Amended) Composite indefinite chill rolls according to claim 26 [through 28], characterized in that the working or sleeve material has, in wt-%,
- V = 3.1 to 3.9, preferably 3.3 to 3.75 and
- Nb+Ta = less than 0.6
- with the remainder being iron and impurities.
30. (Amended) Composite indefinite chill rolls according to [one of claims 26 through 29] claim 26, characterized in that the binding zone between the sleeve or working part and the roll core of low-alloy cast iron, preferably of ductile iron, has, in the radial direction, a bending strength (3-point bending test) of greater than 600 N/mm².

**Casting Material for Indefinite Chill Rolls with a Sleeve Part and Method for
Producing the Same**

The invention relates to a method for the production of alloyed casting material, in particular of material for the working area of indefinite chill rolls, containing the elements carbon, silicon, manganese, chromium, nickel, molybdenum, vanadium, and optionally also other elements of group 5 of the periodic system, aluminum, and the remainder iron and impurities related to the manufacturing process.

The invention further relates to a casting material containing the elements listed above.

Finally, the invention includes a composite indefinite chill roll, especially for work rolls for forming flat steel, comprising a work or sleeve part made of a casting alloy with little tendency to adhere or weld to the rolling stock and of a high-strength core part made of low-alloy cast iron, especially of ductile iron.

Tools or machine parts that are exposed to numerous stresses of different types require a special set of properties. For this reason, the most appropriate materials and manufacturing process must be selected for each case with regard to feasibility and economical manufacture as well as the service life of the parts in practical use.

Parts which are exposed to varying temperatures above room temperature, especially for hot forming of workpieces, are made of casting materials in many applications. As a result of such a choice of material, distortion due to local differences in temperature can advantageously be minimized to a great extent, the manufacture of the parts can be made economical, and the material properties can be largely matched to the stresses.

Work rolls used for hot rolling of steel, for example, rolls in wide-strip hot rolling mills, especially in Steckel stands and in the finishing stands, on the one hand, are

exposed to high mechanical and thermal stresses and, on the other hand, must exhibit the least possible tendency toward sticking and welding to the rolling stock. Specifically, the thinner the hot strip is rolled, the higher the specific contact pressures become between the work roll and rolling stock, which significantly increases a tendency toward adhesion of the strip to the roll surface, enhanced by the low final roll temperatures of the last stands. This sticking or welding of the roll surface to the strip can lead to tearing of material out of the strip, which in and of itself and through its adhesion to the roll can lead to further rolling defects, which often perforce leads to a reduced quality in the hot rolled strip.

It is known to use an indefinite chill cast material in the working area of the rolls in the last stands of a hot strip rolling mill in order to meet the requirements regarding a reduction of friction in the roll gap and reduction of a tendency of the strip to stick to the roll surface, and also to increase the resistance to stripping and thermal shock damage to the material.

The indefinite chill quality comprises three essentially different structural constituents that are formed as cast, specifically graphite, carbides, and a steel-like matrix. Only the matrix can be significantly altered by heat treatment. The indefinite chill roll quality and/or alloy forms a large amount of carbide and little graphite in the structure when solidification is rapid, and at lower solidification rates the relationship is reversed, i.e., less carbide and more graphite is formed. This has the consequence that rapidly solidified material is harder and slowly solidified material is softer. In an indefinite chill roll, this has the effect that the carbide content drops as the distance from the cast surface increases, the graphite content rises, and the hardness likewise decreases. Since no defined jump in hardness can be observed in this case, this quality has been named "indefinite."

However, graphite precipitation can degrade the hardness and especially the wear properties of the material, so the microstructure should also contain hard carbides to minimize this disadvantage.

It is known to one skilled in the art to create a cast structure with graphite particles and carbides using alloying methods, where the content of elements that promote graphite formation, primarily nickel and silicon, and the content of carbide formers, primarily small amounts of chromium and molybdenum, as well as the carbon content in the melt must all be coordinated with one another and where their interactions during solidification must also be taken into account.

According to the prior art, indefinite chill rolls have a composition in wt-% of 2.6 to 3.6 % carbon, 0.6 to 1.1 % silicon, 0.6 to 1.0 % manganese, 1.5 to 2.1 % chromium, 4.1 to 4.6% nickel, 0.3 to 0.5% molybdenum, with the remainder being iron, accompanying elements, and impurities. The structure of the working body or the jacket of a composite roll is primarily composed of a bainitic and/or martensitic matrix with constituents of 28 to 40 % of eutectic carbides and 1.3 to 2.2 % graphite by volume, with 5 to 20 graphite particles being present per mm² of polished surface.

In order to improve the performance characteristics of indefinite chill rolls, especially to increase their wear resistance in the working area, attempts have already been made (PCT/GB 93/02380) to introduce preferable surface-layered carbide particles of greater hardness into the melt provided for this purpose. It is known to practitioners of the art that small amounts of high-hardness carbides increase wear resistance of the material more than the increase produced by typical-quality carbides of lower hardness. If the roll or the roll jacket is made from a melt of this type by means of a centrifugal casting process, undesirable segregation and inhomogeneities can be produced as a result of the centrifugal force and the differences in specific gravity between the carbide particles and the melt. Furthermore, the formation of the necessary graphite can be disrupted by the change in the melt.

The proposal was made according to PCT/US 96/09181 to add 0.3 to 6.0 wt-% Nb to a melt having a balanced composition for indefinite chill rolls and to correspondingly increase the carbon content stoichiometrically with respect to the niobium carbide to be formed. While this process does increase the carbide content and

the wear resistance of the material, higher niobium contents can lead to primary formation of carbides which can result in coarsening of the carbide grains and the graphite particles.

5 During the centrifugal casting of the working area of an indefinite chill roll, the alloy in the mold is subjected to a high centrifugal acceleration during solidification, for example, in the range of 80 to 180 g. Since the monocarbides of vanadium that are primarily formed in the melt have a lower density than the liquid metal, and those of niobium have a higher density, segregation and/or demixing can occur. A proposal has
10 already been made (US 5 738 734) to prevent segregation by alloying the melt equally with vanadium and niobium in such a manner that the monocarbides formed during solidification are mixed carbides (VNb)C and have essentially the same density as the melt. As a result of the highest possible content for the monocarbide-forming elements of 17 wt-% per the above U.S. Patent, the carbon concentration must also be adjusted
15 according to the known relationship. However, such an alloy can exhibit an undesirable solidification structure with localized demixing and large graphite particles, which can produce impaired surface quality of the roll after even a short operating time, and also increases the tendency to stick.

20 Based on the prior art, the object of the invention is to specify a new, improved process by means of which the material of the working area of indefinite chill rolls has a significantly reduced tendency to stick or weld to the rolling stock and a consistently high abrasion resistance over the thickness of the area used.

25 The invention has the additional goal of producing a casting material that has finely dispersed and homogeneously distributed graphite precipitations with a low volume fraction, has special carbides with uniformly small grain diameter likewise uniformly distributed through the base material, and also has essentially unchanged working surface characteristics in the event of wear.

30 Lastly, the invention has achieved the object of producing composite indefinite

chill rolls having significantly improved working characteristics and reduced danger of roll breakage, stripping and cracking in the core transition region.

This object is attained in a generic process in that:

5 A. A melt is produced that has a chemical composition in wt-% of

2.0 to 3.5 C

1.0 to 2.0 Si

0.5 to 2.0 Mn

1.0 to 3.0 Cr

10 3.5 to 4.9 Ni

0.2 to 2.9 Mo

with the remainder iron and impurities, and

B. More than 0.5 wt-% vanadium in amounts up to 5.9 wt-% is added, is dissolved therein, and

15 C. The composition of the melt is set using alloying methods by fixing the concentrations of carbon and silicon in the presence of nickel and the effective sum of the carbide forming elements in such a manner that, at its solidification, a microstructure is formed which exhibits 1.0 to 3.0 vol-% of graphite, with the guideline that more than 20 and less than 100 graphite particles are present per mm² of observed surface in a metallographic section and the remainder is primarily composed of martensite, 8 to 35 vol-% of eutectic carbides, and at least 1 vol-% of finely distributed vanadium carbides, after which

20 D. The melt is cast in a form, preferably a centrifugal casting form, and is allowed to solidify into a body, preferably a working body of a roll, and if necessary the cast body is further processed, for example, into a composite roll, with the body or roll that has been produced in this manner being

25 E. Subjected to heat treatment including at least one one-time heating to a treatment temperature, holding at this temperature, and cooling to room temperature.

30 The advantages achieved by the invention lie essentially in the fact that the solidification kinetics of the melt and the structural morphology of the material have been set so as to be advantageously changed. This change is achieved through the synergistic

effect of the alloying elements in the specified concentrations, where a high number of small graphite particles is made possible by a slight increase, as compared to the prior art, in the silicon content and also aluminum in the presence of nickel in within tight limits. In this context, however, the effective total of the carbide-forming elements during eutectic solidification is important, where chromium and molybdenum in the provided concentrations were found to be critical influencing variables. Because at least some of the vanadium carbides are precipitated prior to eutectic solidification to the solubility limit of vanadium in the liquid alloy, it is important for these monocarbides to have a small particle size and thus to be incapable of segregating in the melt during solidification as a result of the applied centrifugal acceleration. According to the current state of knowledge, the fine-grain nature of the primary dispersed carbide precipitation is achieved through the interaction of carbon, silicon and nickel, on the one hand, and chromium, molybdenum and vanadium, on the other hand. These interactions of the activities of the elements are still not completely understood by science, but it can be assumed that advantageous precipitation kinetics are achieved during solidification and that with appropriate silicon content and nickel concentrations in the remaining melt the precipitation of graphite and eutectic carbides is delayed and that fine-grain residual solidification takes place once greater supercooling is achieved. The composition of the melt here should be set in such a way that the graphite portion in the solidified material is 1.0 to 3.0 vol-%. Lower graphite percentages increase the tendency of the rolling stock to stick to the roll surface even at a high graphite particle density of over 20 per mm². When the graphite content exceeds 3.0 vol-%, roll wear increases. Moreover, a content of between 8 to 35 vol-% of eutectic carbides and a content of at least 1 vol-% of special carbides or monocarbides is to be produced by alloying methods. Carbide proportions below 8 and 1 vol-% lead to reduced wear resistance of the material, and more than 35 vol-% of eutectic carbides increases the risk of cracking or breakage.

Particularly pronounced crazing resistance as well as surface quality with low roll wear in operation can be achieved when the composition of the melt is established by alloying so that a microstructure is formed at solidification which has 1.2 to 2.5 vol-%, preferably 1.25 to 1.95 vol-%, of graphite with the guideline that more than 22 but no

more than 90 graphite particles are present per mm² of observed surface in a metallographic section and the remainder is primarily composed of martensite, 10 to 25 vol-% of eutectic carbides, and 2 to 20 of finely distributed monocarbides.

When, according to a preferred refinement of the invention, the composition of the melt is set such that, in the presence of nickel, the concentration ratio of carbon to silicon is less than or equal to 2.6, preferably less than or equal to 2.0, the graphite precipitation and/or the graphite fraction in the material can be held within the desired range with high precision and within tight limits. At a ratio of carbon content to silicon content that exceeds 2.6, coarse primary carbides are formed and graphite formation is adversely affected as well.

In the optimization of material properties and material quality, although it is advantageous for the carbon content of the melt to be set to a value of 2.2 to 3.1 wt-%, preferably 2.6 to 2.95.

In the interests of an especially good balance of the graphite and carbide distribution during solidification, and in order to further improve the operating characteristics of the roll, it has been proven favorable for a final silicon content of more than 1.2 to 1.95 wt-%, preferably 1.4 to 1.75, to be provided.

The element aluminum fosters the tendency toward graphite formation, on the one hand, but also causes fine-grain precipitation of special carbides, on the other hand. Aluminum can also partially replace silicon in terms of action kinetics and find application as a control element for a balanced graphite/carbide precipitation so that, when setting the composition of the melt by alloying means, aluminum can be added at 0.002 to 0.65 wt-% and dissolved therein. Aluminum contents of 0.005 to 0.04 wt-% are preferred.

The establishment of high material quality in tight limits is favorable when the nickel content of the melt is set to a value of 3.51 to 4.7 wt-%, preferably 4.15 to 4.6 wt-%.

5 %.

In terms of solidification kinetics, but also with regard to the formation of a large number of graphite particles, it has proven advantageous for the concentration ratio of molybdenum to chromium to be less than 1.0, preferably less than 0.8.

$\text{Mo/Cr} < 1.0$, preferable < 0.8

If the ratio exceeds 1.0, high transformation stresses can arise during cooling and heat treatment of the composite roll, which can lead to material separation. This danger is greater with smaller rolls; however, for reasons of safety from cracking it is always advantageous to set the ratio of chromium content to molybdenum content below 0.8.

In the interest of the deliberate formation of eutectic carbides and thus the reduction of the risk of breakage of the roll material under impact stresses, it has proven advantageous for the content of chromium and molybdenum in the melt in wt-% to be set to the values of

chromium 1.2 to 2.6, preferably 1.5 to 2.01
molybdenum 0.20 to 2.6, preferably 0.3 to 0.9.

Manganese serves primarily to bind sulfur, where it is advantageous for the manganese content in the melt to be set to a value in wt-% of 0.6 to 1.6, preferably 0.7 to 1.45.

In order to further promote a finely dispersed distribution of graphite particles, and to keep the grain size of the special carbides uniformly small, and thus improve the working properties of an indefinite chill roll, even under frequent abrasion, it can be of further advantage for 1.8 to 3.9 wt-%, preferably 1.9 to 2.9 wt-%, of vanadium to be added to the melt and dissolved therein.

It can also be advantageous when some of the vanadium is replaced by additional

elements from group 5 of the periodic system in an amount of less than 0.6 wt-%, and mixed carbides are formed. In the end, the prescribed material properties are achieved by means of heat treatment. In the course of the method according to the invention, it has proven advantageous for the cast body or the roll to be subjected to a heat treatment including at least one heating from room temperature to a treatment temperature of 400 °C to 500 °C, preferably 460 °C to 480 °C, holding at this temperature for at least two hours, preferably at least 8 hours, and cooling to room temperature, optionally with a tow-temperature treatment.

The further object of the invention is achieved with a casting material of the initially described type in that the alloy contains, in wt-%,

0.5 to 5.9 V

1.0 to 2.0 Si

0.5 to 2.0 Mn

1.0 to 3.0 Cr

3.5 to 4.9 Ni

0.20 to 2.9 Mo

2.0 to 3.5 carbon with the guideline that

1.0 to 3.0 vol-% is present as graphite particles with a distribution of more than 20 and less than 100 graphite particles per mm² of metallographic polished surface of the material.

The advantage of the material thus formed is its special suitability for construction of indefinite chill rolls, and can largely be observed in that, as compared to the prior art, a very constant graphite proportion is achieved with similar properties of the material. Due to the high graphite particle density brought about by alloying methods, the tendency of the rolling stock to weld or adhere to the surface of the roll is reduced significantly. A particle count below 20 per mm² of observation surface does not demonstrate sufficient effect, however. The same applies when the number of graphite particles is above 100 per mm² because then the diameter of the individual particles is too small to reduce the tendency to adhere to the required degree. For a high graphite particle count and a fine

eutectic solidification with small special carbides, it is necessary for the vanadium content to be greater than 0.5 wt-% because smaller concentrations do not achieve effective structural refinement. In order to achieve formation of finely dispersed graphite particles and to maintain a desired solidification and structural arrangement when the requisite carbon content is present, it is also necessary to provide the elements silicon in the presence of nickel and also chromium and molybdenum within tight concentration limits, because these elements interact kinetically. However, a high vanadium content leads to coarse primary carbide precipitation of the MC type, and can cause increased fracture risk and flaking of the large carbides out of the working surface, so the concentration of this element in the material should not exceed the value of 5.9 wt-%.

The service properties of the roll can be advantageously further improved when the alloy contains

1.8 to 4.8 wt-% of elements of the vanadium group of the periodic system

2.2 to 3.1 carbon with the guideline that

1.2 to 2.5 vol-% of graphite is present as particles with a distribution of more than 22 particles and less than 90 particles per mm² of a metallographically polished surface. If the graphite particle count of 100 per mm² of image surface is exceeded for a graphite content of 1.8 vol-%, the tendency of the rolling stock to stick to the roll surface increases significantly.

High quality assurance, especially with regard to the transformation behavior of the material, is achieved when the alloy contains, in wt-%,

2.0 to 3.5 carbon

1.0 to 2.0 silicon

0.5 to 2.0 manganese

1.0 to 3.0 chromium

3.5 to 4.9 nickel

0.2 to 2.9 molybdenum

1.5 to 4.9 vanadium

with the remainder iron and impurities.

Furthermore, as became evident, it is advantageous in the formation according to the invention of the material composition with regard to a homogeneous and finely dispersed graphite particle formation as well as improved service properties of an indefinite chill roll if the alloy has a concentration ratio of carbon to silicon of less than or equal to 2.6, preferably less than or equal to 2.0, where the presence of nickel is provided.

It has proven advantageous, both for especially fine graphite and carbide formation as well as for balanced eutectic graphite/carbide precipitation, for the alloy to contain greater than 1.2 to 1.85 wt-% of silicon, preferably from 1.4 to 1.75 wt-%.

Aluminum, in content levels from 0.002 to 0.65 wt-%, preferably from 0.005 to 0.04 wt-%, can advantageously ensure a desired graphite and carbide formation as well as a fine solidification structure of the cast body.

With regard to a controlled graphite content and an intended assumption of hardness of the material, it is preferred for the alloy to contain 3.5 to 4.9 wt-% of nickel, preferably 4.15 to 4.6 wt-%.

In order to bind the sulfur, the alloy can beneficially contain 0.6 to 1.6 wt-%, preferably 0.6 to 1.6 wt-%, of manganese.

Both the solidification morphology and the transformation behavior of the sleeve material can be improved, and the risk of cracking of the composite roll can be reduced, when the alloy has a concentration ratio of molybdenum to chromium of less than 1.0, preferably less than 0.8. In this way, the inner stresses of a roll are significantly reduced. This applies to vanadium content levels of up to 5.9 wt-% and only low content levels of further elements from group 5 of the periodic systems. The proportion of eutectic carbides in the roll material can be advantageously developed by content levels in wt-% of 1.5 to 2.01 of chromium and concentrations from 0.3 to 0.9 of molybdenum, especially with a carbon content from 2.6 to 2.95 wt-%.

When the alloy contains 1.8 to 4.0 wt-% of vanadium, preferably 1.9 to 2.95 wt-%, favorable resistance to wear with high material hardness and an improved structural transformation behavior of the material are achieved at the same time.

5 It is also possible to replace some of the vanadium content with other monocarbide-forming elements from group 5 of the periodic system in amounts less than 0.6 wt-%. At niobium or tantalum concentrations in the alloy of 0.6 wt-% and greater, coarse phases may be formed in the structure which worsen the properties of the working roll and the surface quality of the rolling stock.

10 Finally, improved resistance to breakage and a low stripping tendency with improved wear characteristics of the cast material can be achieved if the cast material possesses, in vol-%,
8 to 35, preferably 10 to 25, eutectic carbide and
15 1 to 15, preferably 2 to 10, carbides from group 5, the vanadium group, of the periodic system.

20 The further object of the invention, namely to specify composite indefinite chill rolls of the generic type produced with the centrifugal casting process and having significantly improved working characteristics and reduced danger of roll breakage, stripping, crazing and cracking in the transition region to the core, is achieved in that the working area or sleeve has a thickness of 10 to 150 mm, and the sleeve material has a structure comprising primarily of 1.0 to 2.5 vol-% of graphite, where the latter is present in finely dispersed form with a graphite particle count of more than 20 particles per mm²
25 of a metallographic polished surface, and of 8 to 35 vol-% of eutectic carbides, of 1 to 20 vol-% of vanadium carbides that are uniformly distributed, especially in the direction of the sleeve thickness, with the remainder being essentially martensite and constituents related to impurities or the manufacturing process, and has a hardness between 70 and 90 ShC.

30 The advantage of the rolls according to the invention is essentially that the sleeve

metallically bound to the core with high strength has a high graphite particle count, which particularly effectively prevents adhesion or welding of the rolling stock during roll operation. This homogeneous graphite formation and the uniform distribution of the small vanadium special carbides is achieved by using alloying methods to influence the solidification kinetics so that demixing due to so-called centrifugal segregation cannot take place during the centrifugal casting process. Thus, in an advantageous fashion, the structural arrangement and the roll performance are largely the same after every instance of reworking the working surface, even in the event of necessary radial grinding. The roll performance in each case up until a necessary reworking of the surface is advantageously improved because the high graphite particle density effects an increased crazing resistance as well as an improved surface quality of the sleeve, which is more wear-resistant as a result of the special carbides.

Improved properties of a roll according to the invention can be attained with certainty when the working area or sleeve material has a structure which contains 1.0 to 2.5 vol-% of graphite, with the guideline that its distribution density is at least 22 particles but less than 100 graphite particles per mm^2 of polished surface in a metallographic section, contains eutectic carbides in an amount of 10 to 25 vol-%, and 2 to 10 vol-% of special carbides of the elements of group 5 of the periodic system

If, according to a preferred material variant, the working or sleeve material has a composition, in wt-%, of

C = 2.0 to 3.5, preferably 2.21 to 3.1, especially 2.6 to 2.95

Si = 1.0 to 2.0, preferably over 1.2 to 1.85, especially 1.4 to 1.75

Mn = 0.5 to 2.0, preferably 0.6 to 1.6, especially 0.7 to 1.4

Cr = 1.0 to 3.0, preferably 1.3 to 2.5, especially 1.5 to 2.01

Ni = 3.5 to 4.9, preferably 3.5 to 4.7, especially 4.15 to 4.6

Mo = 0.2 to 2.9, preferably 0.25 to 1.3, especially 0.3 to 0.9

Al = 0.002 to 0.65, preferably 0.005 to 0.1, especially 0.005 to 0.04

V = 0.5 to 5.9, preferably 1.8 to 3.9, especially 1.9 to 2.9

optionally, Nb and/or Ta less than 0.6

with the remainder being iron and impurities and the roll core is formed of ductile iron, then high resistance to wear, reduced danger of crack formation and crack propagation, and high hardness of the working area of the roll are assured.

High protection from the initiation of cracks can be achieved when the binding zone between the sleeve or working area and the roll core of low-alloy cast iron, preferably of ductile iron, has, in the radial direction, a bending strength (3-point bending test) of greater than 600 N/mm^2 .

The invention is described in detail below using graphs and illustrations of test results as well as a table. Shown are:

Fig. 1 a graph of C/Si

Fig. 2 a graph of Mo/Cr

Figs. 3 and 4 unetched photomicrographs

Tab. 1 roll materials and their performance in practical use.

Fig. 1 shows the concentration of silicon and carbon, where the inventive region is described by the points α , β , γ , δ . Preferred regions with a ratio of $\text{C/Si} = 2.6$ (region A) (α , β , γ , δ^1 , α^1) and a ratio of $\text{C/Si} \leq 2.0$ (region B) (α , β , γ , δ^2) are marked.

Fig. 2 shows a graph of molybdenum and chromium in which the ratio region (α , β , γ , δ) according to the invention of the content is represented. The preferred regions with a ratio of $\text{Mo/Cr} \leq 1.0$ (region A) (α , β , γ , δ , δ^1 , α^2) and one with $\text{Mo/Cr} = 0.8$ (region B) (α , β , γ , δ , δ^2 , α^1) are identified as in Fig. 1.

Fig. 3 shows, in a photomicrograph with 50x magnification, the formation of

graphite in a roll material according to the prior art. The roll sleeve had the following chemical composition in wt-%: C = 3.09, Si = 0.91, Mn = 0.84, Cr = 1.79, Ni = 4.51, Mo = 0.38, Al = 0.003, graphite content: 3.9 vol-%, 18 graphite particles per mm². Fig. 4 shows, in a photomicrograph with the same 50x magnification, the large number and uniform distribution of graphite particles in a working area of a roll composed according to the invention. The chemical composition of the working area in wt-% was C = 3.02, Si = 1.42, Mn = 0.9, Cr = 1.8, Ni = 4.36, Mo = 0.52, V = 2.9, Al = .008, graphite content: 2.8 vol-%, 42 graphite particles per mm².

In comparison to a material according to the prior art, despite a lower carbon content and lower graphite content of the alloy according to the invention, the number of graphite particles was more than twice as large and 3.2 vol-% of vanadium carbides were measured.

Table 1 summarizes the chemical composition of the roll sleeve, the structural composition and the roll performance achieved in practical use for each of 10 roll pairs. The rolls designated A through E, which were manufactured according to prior art, hence were not alloyed with vanadium; the rolls designated F through J were manufactured with a sleeve material alloyed according to the invention.

By means of additional alloying with vanadium (rolls F through N), the formation of hard vanadium carbides with small grain size and largely homogeneous distribution in the material was possible with a reduced proportion of eutectic carbides, by means of which the wear resistance of the material, and finally the roll performance, were significantly improved. A high graphite particle count per mm², which was achieved through the interactions of the activities of the elements Cr, Si, Ni, Mo, C and V, prevented sticking or welding of the rolling stock to or with the roll surface, even at low graphite proportions. The addition of niobium and tantalum, which is to say of additional elements from group 5 of the periodic system, produced a small increase in the wear resistance and roll performance in operation at contents less than 0.6 wt-%. It is noteworthy that the formation and propagation of cracks, as well as stripping, were

significantly reduced in the sleeve material according to the invention, which can probably be attributed to the large number of graphite particles. Microscopic examination demonstrated that the monocarbides MC had small grain sizes and were primarily distributed in finely dispersed form. Since the density of the vanadium carbides is approximately 5.82 g/cm³ at room temperature, on the one hand, and no centrifugal segregation caused by the centrifugal casting was observable on the other hand, we can conclude that the precipitation of special carbides and the fine precipitation of graphite occurred primarily during the eutectic solidification and that primary precipitation was largely inhibited.

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Des.	Composition of the sleeve										Structural Composition								Roll performance
	C wt-%	Si wt-%	Mn wt-%	Cr wt-%	Ni wt-%	Mo wt-%	V wt-%	Nb+Ta wt-%	Al wt-%	C/Si	Mo/Cr	carbides, total vol-%	carbides, eutectic vol-%	carbides of group 5 vol-%	graphite portion vol-%	graphite particles per mm ²	hardness ShC	t/mm	
5	A	3.34	0.81	0.94	1.74	4.31	0.38	-	-	0.002	4.12	0.22	33.2	33.2	0	3.2	46	75-78	3.183
	B	3.27	0.84	1.04	1.73	4.3	0.38	-	-	0.002	3.89	0.22	35.1	35.1	0	2.6	19	76-80	3.868
	C	3.11	0.91	0.84	1.79	4.51	0.38	-	-	0.003	3.42	0.21	31	31	0	2.8	21	77-79	2.417
	D	3.09	0.91	0.81	1.71	4.52	0.38	-	-	0.004	3.40	0.22	28.7	28.7	0	3.9	18	77-78	2.915
	E	3.32	1.02	0.78	1.75	4.4	0.4	-	-	0.002	3.25	0.23	29.5	29.5	0	5.3	62	77-79	1.736
10	F	2.75	1.42	0.9	1.8	4.36	0.85	2.9	-	0.008	1.94	0.47	27.3	24.1	3.2	1.7	42	76-80	6.253
	G	2.83	1.45	0.89	1.79	4.37	0.82	2.8	-	0.008	1.95	0.46	25.8	22.9	2.9	1.9	38	76-79	6.253
	H	3.05	1.43	0.92	1.82	4.45	1.2	2.83	-	0.011	2.13	0.66	26	22.7	3.3	2.8	37	76-79	5.434
	I	2.9	1.65	0.93	1.93	4.27	0.85	3.35	0.52	0.006	1.76	0.44	21.3	14	7.3	1.8	24	80-83	3.503 *)
	J	2.93	1.71	0.95	1.85	4.28	0.35	2.75	0.35	0.012	1.71	0.19	18.7	12.7	6	1.7	33	76-79	6.867
15	K	2.9	1.52	0.92	1.62	4.32	1.53	3.24	-	0.009	1.91	0.94	21.4	17.2	4.2	2.3	40	77-80	6.407
	L	2.83	1.59	1.02	1.87	4.2	0.85	2.85	-	0.015	1.78	0.45	21.4	18.2	3.2	1.8	27	76-78	6.684
	M	2.91	1.6	0.85	1.94	4.15	1.42	3.25	0.27	0.017	1.82	0.73	21.6	16.5	5.1	1.9	53	80-83	6.173
	N	2.87	1.53	0.97	1.72	4.27	1.53	3.14	0.45	0.021	1.88	0.89	25.7	20.4	5.3	2.2	35	79-81	6.290

*) Particularly difficult roll conditions

F, G, H, I, J, K, L, M, N: rolls according to the invention

Table 1

Claims

1. Method for the production and processing of alloyed casting material for the working area of indefinite chill rolls, containing the elements carbon, silicon, manganese, chromium, nickel, molybdenum, vanadium, and if applicable additional elements of group 5 of the periodic system, aluminum, and the remainder iron, accompanying elements, and impurities related to the manufacturing process, characterized in that

A. a melt is produced that has a chemical composition in wt-% of

2.0 to 3.5 C

1.0 to 2.0 Si

0.5 to 2.0 Mn

1.0 to 3.0 Cr

3.5 to 4.9 Ni

0.2 to 2.9 Mo

with the remainder iron and impurities, and

B. more than 0.5 % vanadium by weight in amounts up to 5.9 wt-% is added, is dissolved therein, and

C. the composition of the melt is set using alloying methods by fixing the concentrations of carbon and silicon in the presence of nickel and the effective total of the carbide forming elements in such a manner that, at its solidification, a microstructure is formed which exhibits 1.0 to 3.0 vol-% of graphite, with the guideline that more than 20 but less than 100 graphite particles are present per mm² of observed surface in a metallographic section and the remainder is composed primarily of martensite, 8 to 35 vol-% of eutectic carbides, and at least 1 vol-% of finely distributed vanadium carbides, after which

D. the melt is cast in a form, preferably a centrifugal casting mold, and is allowed to solidify into a body, preferably a working body of a roll, and optionally, the cast body is further processed, for example, into a composite roll, with the body or roll that has been produced in this manner being

E. subjected to a heat treatment including at least one one-time heating to a

treatment temperature, holding at this temperature, and cooling to room temperature.

2. Method according to claim 1, characterized in that the composition of the melt is set using alloying methods by fixing the concentrations of carbon and silicon in the presence of nickel and the effective total of the carbide forming elements in such a manner that, at its solidification, a microstructure is formed which has 1.0 to 2.5 vol-% of graphite, with the guideline that more than 22 and less than 100 graphite particles are present per mm² of observed surface in a metallographic section and the remainder is composed primarily of martensite, 10 to 25 vol-% of eutectic carbides, and of 2 to 20% of finely distributed carbides of the elements of group 5 of the periodic system.

3. Method according to one of claims 1 or 2, characterized in that the composition of the melt is set in such a manner that the concentration ratio of carbon to silicon is less than or equal to 2.6, preferably less than or equal to 2.0:

$$C/Si \leq 2.6, \text{ preferably } = 2.0$$

4. Method according to one of claims 1 through 3, characterized in that the carbon content of the melt is set to a value of 2.2 to 3.1 wt-%, preferably 2.6 to 2.95%.
5. Method according to one of claims 1 through 4, characterized in that a final content of silicon of 1.2 to 1.85 wt-%, preferably 1.4 to 1.75%, is provided.
6. Method according to one of claims 1 through 5, characterized in that, when the composition of the melt is set using alloying methods, 0.002 to 0.65 wt-%, preferably 0.005 to 0.04 %, of aluminum is added and dissolved therein.
7. Method according to one of claims 1 through 7, characterized in that the nickel content of the melt is set to a value of 3.51 to 4.7 wt-%, preferably 4.15 to 4.6 wt-%.

%.

8. Method according to one of claims 1 through 7, characterized in that the composition of the melt is set in such a manner that the concentration ratio of molybdenum to chromium is less than 1.0, preferably less than 0.8:

$\text{Mo/Cr} < 1.0$, preferably < 0.8 .

9. Method according to one of claims 1 through 8, characterized in that the content levels of chromium and molybdenum in wt-% of the melt are set to the values of chromium 1.5 to 1.9 molybdenum 0.3 to 0.9.

10. Method according to one of claims 1 through 9, characterized in that 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.9 wt-%, is added to the melt and dissolved therein.

11. Method according to one of claims 1 through 10, characterized in that some of the vanadium is replaced by additional elements from group 5 of the periodic system in an amount of less than 0.6 wt-%, and mixed carbides are formed.

12. Method according to one of claims 1 through 11, characterized in that the cast body or the roll is subjected to a heat treatment which comprises heating from room temperature to a treatment temperature of 400 °C to 500 °C, preferably 460 °C to 480 °C, holding at this temperature for at least two hours, preferably at least 8 hours, and cooling to room temperature, optionally with a low-temperature treatment.

13. Casting material for the working area of indefinite chill rolls, containing the elements carbon, silicon, manganese, chromium, nickel, molybdenum, vanadium, and the remainder iron, accompanying elements, and impurities related to the

manufacturing process, characterized in that the alloy contains, in wt-%, more than

0.5 to 5.9 vanadium

1.0 to 2.0 silicon

0.5 to 2.0 manganese

1.0 to 3.0 chromium

3.5 to 4.9 nickel

0.20 to 2.9 molybdenum

2.0 to 3.5 carbon with the guideline that

1.0 to 3.0 vol-% is present as graphite in particles with a distribution of more than 20 and less than 100 particles per mm² of polished surface of the material.

14. Casting material according to claim 13, characterized in that the alloy contains

1.8 to 4.9 wt-% vanadium

2.2 to 3.1 wt-% carbon with the guideline that

1.2 to 2.5 vol-% as graphite in particles with a distribution of more than 22 and less than 90 particles per mm² of polished surface.

15. Casting material according to claim 13 or 14, characterized in that the alloy contains, in wt-%,

2.0 to 3.5 carbon

1.0 to 2.0 silicon

0.5 to 2.0 manganese

1.2 to 2.5 chromium

3.5 to 4.9 nickel

0.5 to 2.1 molybdenum

1.5 to 4.9 vanadium

with the remainder iron and impurities.

16. Casting material according to one of claims 13 through 15, characterized in that

the alloy has a concentration ratio of carbon to silicon of less than or equal to 2.6, preferably less than or equal to 2.0:

$C/Si \leq 2.6$, preferably ≤ 2.0

5

17. Casting material according to one of claims 13 through 16, characterized in that the alloy contains 2.6 to 2.95% carbon by weight.

18. Casting material according to one of claims 13 through 17, characterized in that the alloy contains 1.2 to 1.85 wt-% of silicon, preferably 1.4 to 1.75%

10

19. Casting material according to one of claims 13 through 18, characterized in that the alloy contains 0.002 to 0.65 wt-% of aluminum, preferably 0.005 to 0.04 %.

15

20. Casting material according to one of claims 13 through 19, characterized in that the alloy contains 3.5 to 4.9 wt-% of nickel, preferably 4.15 to 4.6%.

21. Casting material according to one of claims 13 through 20, characterized in that the alloy has a concentration ratio of molybdenum to chromium of less than 1.0, preferably less than 0.8:

20

$Mo/Cr < 1.0$, preferably < 0.8 .

22. Casting material according to one of claims 13 through 21, characterized in that the alloy contains in wt-%
chromium 1.5 to 2.01
molybdenum 0.3 to 0.9.

25

23. Casting material according to one of claims 13 through 22, characterized in that the alloy contains 1.8 to 3.9 wt-% of vanadium, preferably 1.9 to 2.95 wt-%.

30

24. Casting material according to one of claims 13 through 23, characterized in that some of the vanadium content is replaced by additional elements from group 5 of the periodic system in a proportion of less than 0.6 wt-%.

25. Casting material according to one of claims 13 through 24, characterized in that the material possesses, in vol-%, 8 to 35, preferably 10 to 25, eutectic carbides, and 1 to 15, preferably 2 to 10, carbides of the elements of group 5 of the periodic system.

26. Composite indefinite chill rolls, especially for finishing stands of wide strip rolling mills and also Steckel and heavy plate mills, preferably produced using a method according to claims 1 through 12, comprising a work or sleeve part made of a casting alloy, preferably according to claims 13 through 25, with little tendency to adhere or weld to the rolling stock and with a high-strength core part made of ductile iron, characterized in that the working area or sleeve has a thickness of 10 to 150 mm and the sleeve material a structure composed essentially of 1.0 to 2.5 vol-% of graphite, with the latter being finely dispersed with a graphite particle count of more than 20 particles per mm² in a metallographic section, and of 8 to 35 vol-% of eutectic carbides, and of 1 to 20 vol-% of uniformly distributed vanadium carbide, with the remainder composed primarily of martensite and constituents related to impurities or the manufacturing process, and has a hardness between 70 and 90 ShC.

27. Composite indefinite chill roll according to claim 26, characterized in that the working area or sleeve material has a structure which has 1.0 to 2.5 vol-% of graphite, with the guideline that its distribution density is at least 22 particles but less than 100 graphite particles per mm² of polished surface in a metallographic section, contains eutectic carbides in an amount of 10 to 25 vol-%, and 2 to 10 vol-% special carbides of the elements of group 5 of the periodic system.

28. Composite indefinite chill rolls according to claim 26 or 27, characterized in that the working or sleeve material has a composition, in wt-%, of

C = 2.0 to 3.5, preferably 2.21 to 3.1, especially 2.6 to 2.95

Si = 1.0 to 2.0, preferably over 1.2 to 1.85, especially 1.4 to 1.75

Mn = 0.5 to 2.0, preferably 0.6 to 1.6, especially 0.7 to 1.4

Cr = 1.0 to 3.0, especially 1.5 to 2.01

Ni = 3.5 to 4.9, preferably 3.5 to 4.7, especially 4.15 to 4.6

Mo = 0.20 to 2.9, especially 0.3 to 0.9

Al = 0.002 to 0.65, preferably 0.005 to 0.1, especially 0.005 to 0.04

V = 0.5 to 5.9, preferably 1.8 to 3.9, especially 1.9 to 2.9

with the remainder being iron and impurities and the roll core is formed of ductile iron.

29. Composite indefinite chill rolls according to claim 26 through 28, characterized in that the working or sleeve material has, in wt-%,

V = 3.1 to 3.9, preferably 3.3 to 3.75 and

Nb+Ta = less than 0.6

with the remainder being iron and impurities.

30. Composite indefinite chill rolls according to one of claims 26 through 29, characterized in that the binding zone between the sleeve or working part and the roll core of low-alloy cast iron, preferably of ductile iron, has, in the radial direction, a bending strength (3-point bending test) of greater than 600 N/mm².

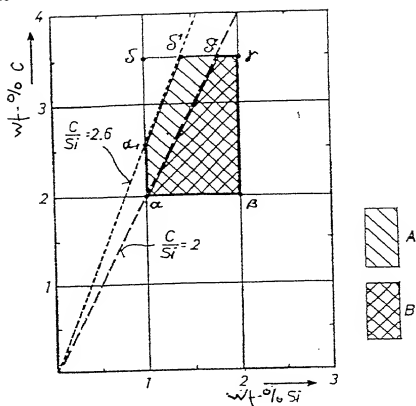


Fig. 1

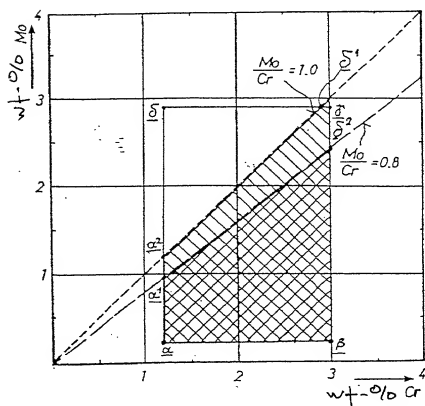
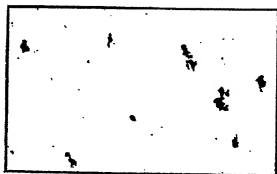


Fig. 2

09/269319

Fig. 3

50 ×

Fig. 4

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German Language Utility or Design Patent Application Declaration

Ich beanspruche hiermit Prioritätsvorteile unter Title 35, US-Code, § 119(e) aller US-Hilfsanmeldungen wie unten aufgezählt.

(Application Number)
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(Status)
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Declaration and Power of Attorney For Utility or Design Patent Application

Erklärung für Patentanmeldungen zur Gebrauchseignung und Entwicklung
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German Language Declaration

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☒ wurde angemeldet am 20 April 2000
unter der US-Anmeldenummer 09/869,310
und wurde am _____ abgeändert (falls zutreffend)
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PCT/AT00/00102
und wurde am 20/Februar/2001 abgeändert (falls
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Ich bestätige hiermit, daß ich den Inhalt der oben angegebene Patentanmeldung, einschließlich der Ansprüche, die eventuell durch einen oben erwähnten Zusatzantrag abgeändert wurde, durchgesehen und verstanden habe.

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As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

CASTING MATERIAL FOR UNDEFINITE ROLLERS WITH A SLEEVE PART AND METHOD FOR PRODUCING THE SAME

the specification of which is attached hereto unless the following box is checked:

☒ was filed on April 20, 2000 as
United States Application Number 09/869,310
and was amended on _____ (if applicable)
or,

PCT International Application Number PCT/AT00/00102
and was amended on February 20, 2001 (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56.

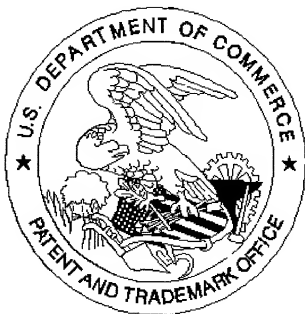
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Priority Claimed
Prioritätsanspruch

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Yes	No
Ja	Nein
<input type="checkbox"/>	<input type="checkbox"/>
Yes	No
Ja	Nein

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